DISEASE:
Craniometaphyseal dysplasia

NAME:	Craniometaphyseal dysplasia
DESCRIPTION:	Craniometaphyseal dysplasia (CMD) is a very rare genetic bone disease characterized by progressive diffuse hyperostosis of cranial bones causing facial dysmorphism and functional repercussions, and metaphyseal widening of long bones.
ORPHACODE:	1522
XREF(S):	Orphanet ICD-10 OMIM OMIM
ANALYTE(S):	ANKH GJA1
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Source URL: http://gentest.healthdata.be/disease/760